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(21) International Application Number: PCT/US94/08581		(74) Agents: KANAGY, James, M. et al.; SmithKline Beecham Corporation, Corporate Intellectual Property, UW2220, 709 Swedeland Road, P.O. Box 1539, King of Prussia, PA 19406-0939 (US).	
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(71) Applicant (for all designated States except US): SMITHKLINE BEECHAM CORPORATION [US/US]; Corporate Intellectual Property, UW2220, 709 Swedeland Road, P.O. Box 1539, King of Prussia, PA 19406-0939 (US).			
(72) Inventors; and (75) Inventors/Applicants (for US only): CHRISTENSEN, Siegfried, B., IV [US/US]; 2216 Race Street, Philadelphia, PA 19103 (US). FORSTER, Cornelia, Jutta [US/US]; 2605 Windsor Drive, Bensalem, PA 19020 (US).			
(54) Title: 3-CYANO-3-(3,4-DISUBSTITUTED)PHENYLCYCLOHEXYL-1-CARBOXYLATES			
(57) Abstract <p>Novel 3-cyano-3-(3,4-disubstituted)phenylcyclohexyl-1-carboxylates or derivatives thereof and their corresponding cyclohexenyl analogs are described herein. These compounds inhibit the production of Tumor Necrosis Factor and are useful in the treatment of disease states mediated or exacerbated by TNF production. These compounds are also useful mediating or inhibiting the enzymatic or catalytic activity of phosphodiesterase IV and are therefore useful in treating diseases in need of mediation or inhibition thereof.</p>			

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3-cyano-3-(3,4-disubstituted)phenylcyclohexyl-1-carboxylates

Field of Invention

The present invention relates to certain novel 3-cyano-3-(3,4-disubstituted)phenylcyclohexyl-1-carboxylates and their corresponding cyclohexenyl
5 analogs, pharmaceutical compositions containing these compounds, and their use in treating allergic and inflammatory diseases and for inhibiting the production of Tumor Necrosis Factor (TNF).

Background of the Invention

Bronchial asthma is a complex, multifactorial disease characterized by reversible
10 narrowing of the airway and hyperreactivity of the respiratory tract to external stimuli.

Identification of novel therapeutic agents for asthma is made difficult by the fact that multiple mediators are responsible for the development of the disease. Thus, it seems unlikely that eliminating the effects of a single mediator will have a substantial effect on all three components of chronic asthma. An alternative to the "mediator approach" is to
15 regulate the activity of the cells responsible for the pathophysiology of the disease.

One such way is by elevating levels of cAMP (adenosine cyclic 3',5'-monophosphate). Cyclic AMP has been shown to be a second messenger mediating the biologic responses to a wide range of hormones, neurotransmitters and drugs; [Krebs
Endocrinology Proceedings of the 4th International Congress Excerpta Medica, 17-29,
20 1973]. When the appropriate agonist binds to specific cell surface receptors, adenylate cyclase is activated, which converts Mg^{+2} -ATP to cAMP at an accelerated rate.

Cyclic AMP modulates the activity of most, if not all, of the cells that contribute to the pathophysiology of extrinsic (allergic) asthma. As such, an elevation of cAMP would produce beneficial effects including: 1) airway smooth muscle relaxation, 2) inhibition of
25 mast cell mediator release, 3) suppression of neutrophil degranulation, 4) inhibition of basophil degranulation, and 5) inhibition of monocyte and macrophage activation. Hence, compounds that activate adenylate cyclase or inhibit phosphodiesterase should be effective in suppressing the inappropriate activation of airway smooth muscle and a wide variety of inflammatory cells. The principal cellular mechanism for the inactivation of cAMP is
30 hydrolysis of the 3'-phosphodiester bond by one or more of a family of isozymes referred to as cyclic nucleotide phosphodiesterases (PDEs).

It has now been shown that a distinct cyclic nucleotide phosphodiesterase (PDE) isozyme, PDE IV, is responsible for cAMP breakdown in airway smooth muscle and inflammatory cells. [Torphy, "Phosphodiesterase Isozymes: Potential Targets for Novel
35 Anti-asthmatic Agents" in New Drugs for Asthma, Barnes, ed. IBC Technical Services Ltd., 1989]. Research indicates that inhibition of this enzyme not only produces airway smooth muscle relaxation, but also suppresses degranulation of mast cells, basophils and neutrophils

along with inhibiting the activation of monocytes and neutrophils. Moreover, the beneficial effects of PDE IV inhibitors are markedly potentiated when adenylate cyclase activity of target cells is elevated by appropriate hormones or autocoids, as would be the case *in vivo*. Thus PDE IV inhibitors would be effective in the asthmatic lung, where levels of
5 prostaglandin E₂ and prostacyclin (activators of adenylate cyclase) are elevated. Such compounds would offer a unique approach toward the pharmacotherapy of bronchial asthma and possess significant therapeutic advantages over agents currently on the market.

The compounds of this invention also inhibit the production of Tumor Necrosis Factor (TNF), a serum glycoprotein. Excessive or unregulated TNF production has been
10 implicated in mediating or exacerbating a number of diseases including rheumatoid arthritis, rheumatoid spondylitis, osteoarthritis, gouty arthritis and other arthritic conditions; sepsis, septic shock, endotoxic shock, gram negative sepsis, toxic shock syndrome, adult respiratory distress syndrome, cerebral malaria, chronic pulmonary inflammatory disease, silicosis, pulmonary sarcoidosis, bone resorption diseases, reperfusion injury, graft vs. host
15 reaction, allograft rejections, fever and myalgias due to infection, such as influenza, cachexia secondary to infection or malignancy, cachexia secondary to human acquired immune deficiency syndrome (AIDS), AIDS, ARC (AIDS related complex), keloid formation, scar tissue formation, Crohn's disease, ulcerative colitis, or pyresis, in addition to a number of autoimmune diseases, such as multiple sclerosis, autoimmune diabetes and systemic lupus
20 erythematosus.

AIDS results from the infection of T lymphocytes with Human Immunodeficiency Virus (HIV). At least three types or strains of HIV have been identified, i.e., HIV-1, HIV-2 and HIV-3. As a consequence of HIV infection, T-cell-mediated immunity is impaired and infected individuals manifest severe opportunistic infections and/or unusual neoplasms. HIV
25 entry into the T lymphocyte requires T lymphocyte activation. Viruses such as HIV-1 or HIV-2 infect T lymphocytes after T cell activation and such virus protein expression and/or replication is mediated or maintained by such T cell activation. Once an activated T lymphocyte is infected with HIV, the T lymphocyte must continue to be maintained in an activated state to permit HIV gene expression and/or HIV replication.

Cytokines, specifically TNF, are implicated in activated T-cell-mediated HIV protein expression and/or virus replication by playing a role in maintaining T lymphocyte activation. Therefore, interference with cytokine activity such as by inhibition of cytokine production, notably TNF, in an HIV-infected individual aids in limiting the maintenance of T cell
30 activation, thereby reducing the progression of HIV infectivity to previously uninfected cells which results in a slowing or elimination of the progression of immune dysfunction caused by HIV infection. Monocytes, macrophages, and related cells, such as kupffer and glial cells, have also been implicated in maintenance of the HIV infection. These cells, like T
35 cells, are targets for viral replication and the level of viral replication is dependent upon the

activation state of the cells. [See Rosenberg *et al.*, The Immunopathogenesis of HIV Infection, Advances in Immunology, Vol. 57, 1989]. Monokines, such as TNF, have been shown to activate HIV replication in monocytes and/or macrophages [See Poli *et al.*, Proc. Natl. Acad. Sci., 87:782-784, 1990], therefore, inhibition of monokine production or activity aids in limiting HIV progression as stated above for T cells.

TNF has also been implicated in various roles with other viral infections, such as the cytomegalovirus (CMV), influenza virus, adenovirus, and the herpes virus for similar reasons as those noted.

TNF is also associated with yeast and fungal infections. Specifically *Candida albicans* has been shown to induce TNF production *in vitro* in human monocytes and natural killer cells. [See Riipi *et al.*, Infection and Immunity, 58(9):2750-54, 1990; and Jafari *et al.*, Journal of Infectious Diseases, 164:389-95, 1991. See also Wasan *et al.*, Antimicrobial Agents and Chemotherapy, 35,(10):2046-48, 1991; and Luke *et al.*, Journal of Infectious Diseases, 162:211-214,1990].

The ability to control the adverse effects of TNF is furthered by the use of the compounds which inhibit TNF in mammals who are in need of such use. There remains a need for compounds which are useful in treating TNF-mediated disease states which are exacerbated or caused by the excessive and/or unregulated production of TNF.

Summary of the Invention

This invention relates to the novel compounds of Formula (I), as shown below, useful in the mediation or inhibition of the enzymatic activity (or catalytic activity) of phosphodiesterase IV (PDE IV). The novel compounds of Formula (I) also have Tumor Necrosis Factor (TNF) inhibitory activity.

This invention also relates to the pharmaceutical compositions comprising a compound of Formula (I) and a pharmaceutically acceptable carrier or diluent.

The invention also relates to a method of mediation or inhibition of the enzymatic activity (or catalytic activity) of PDE IV in mammals, including humans, which comprises administering to a mammal in need thereof an effective amount of a compound of Formula (I), as shown below.

The invention further provides a method for the treatment of allergic and inflammatory disease which comprises administering to a mammal, including humans, in need thereof, an effective amount of a compound of Formula (I).

The invention also provides a method for the treatment of asthma which comprises administering to a mammal, including humans, in need thereof, an effective amount of a compound of Formula (I).

This invention also relates to a method of inhibiting TNF production in a mammal, including humans, which method comprises administering to a mammal in need of such

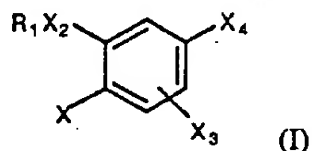
treatment, an effective TNF inhibiting amount of a compound of Formula (I). This method may be used for the prophylactic treatment or prevention of certain TNF mediated disease states amenable thereto.

This invention also relates to a method of treating a human afflicted with a human immunodeficiency virus (HIV), which comprises administering to such human an effective TNF inhibiting amount of a compound of Formula (I).

The compounds of Formula (I) are also useful in the treatment of additional viral infections, where such viruses are sensitive to upregulation by TNF or will elicit TNF production *in vivo*.

The compounds of Formula (I) are also useful in the treatment of yeast and fungal infections, where such yeast and fungi are sensitive to upregulation by TNF or will elicit TNF production *in vivo*.

The compounds of this invention are represented by Formula (I):



wherein:

R_1 is $-(CR_4R_5)_nC(O)O(CR_4R_5)_mR_6$, $-(CR_4R_5)_nC(O)NR_4(CR_4R_5)_mR_6$, $-(CR_4R_5)_nO(CR_4R_5)_mR_6$, or $-(CR_4R_5)_rR_6$ wherein the alkyl moieties may be optionally substituted with one or more halogens;

m is 0 to 2;

n is 1 to 4;

r is 0 to 6;

R_4 and R_5 are independently selected from hydrogen or a C_{1-2} alkyl;

R_6 is hydrogen, methyl, hydroxyl, aryl, halo substituted aryl, aryloxy C_{1-3} alkyl, halo substituted aryloxy C_{1-3} alkyl, indanyl, indenyl, C_{7-11} polycycloalkyl, tetrahydrofuranyl, furanyl, tetrahydropyranyl, pyranal, tetrahydrothienyl, thienyl, tetrahydrothiopyranyl, thiopyranyl, C_{3-6} cycloalkyl, or a C_{4-6} cycloalkyl containing one or two unsaturated bonds, wherein the cycloalkyl and heterocyclic moieties may be optionally substituted by 1 to 3 methyl groups or one ethyl group;

provided that:

a) when R_6 is hydroxyl, then m is 2; or

b) when R_6 is hydroxyl, then r is 2 to 6; or

c) when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then m is 1 or 2; or

d) when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranyl, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then r is 1 to 6;

e) when n is 1 and m is 0, then R₆ is other than H in

-(CR₄R₅)_nO(CR₄R₅)_mR₆;

X is YR₂, halogen, nitro, NR₄R₅, or formyl amine;

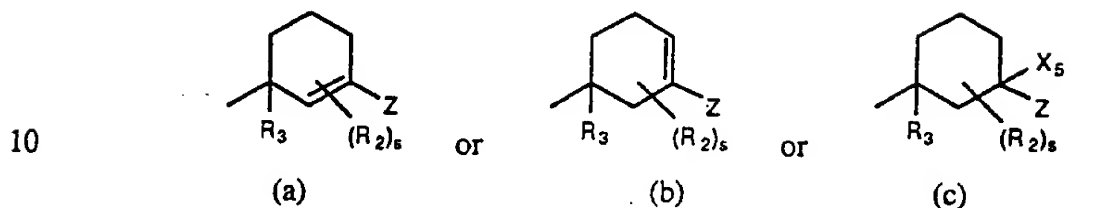
Y is O or S(O)_{m'};

5 m' is 0, 1, or 2;

X₂ is O or NR₈;

X₃ is hydrogen or X;

X₄ is



X₅ is H, R₉, OR₈, CN, C(O)R₈, C(O)OR₈, C(O)NR₈R₈, or NR₈R₈;

15 R₂ is independently -CH₃ or -CH₂CH₃ optionally substituted by 1 or more halogens;

s is 0 to 4;

R₃ is hydrogen, halogen, C₁₋₄ alkyl, CH₂NHC(O)C(O)NH₂, halo-substituted C₁₋₄ alkyl, -CH=CR₈'R₈', cyclopropyl optionally substituted by R₈', CN, OR₈, CH₂OR₈, NR₈R₁₀, CH₂NR₈R₁₀, C(Z')H, C(O)OR₈, C(O)NR₈R₁₀, or C≡CR₈';

20 Z' is O, NR₉, NOR₈, NCN, C(-CN)₂, CR₈CN, CR₈NO₂, CR₈C(O)OR₈, CR₈C(O)NR₈R₈, C(-CN)NO₂, C(-CN)C(O)OR₉, or C(-CN)C(O)NR₈R₈;

25 Z is C(Y')R₁₄, C(O)OR₁₄, C(Y')NR₁₀R₁₄, C(NR₁₀)NR₁₀R₁₄, CN, C(NOR₈)R₁₄, C(O)NR₈NR₈C(O)R₈, C(O)NR₈NR₁₀R₁₄, C(NOR₁₄)R₈, C(NR₈)NR₁₀R₁₄, C(NR₁₄)NR₈R₈, C(NCN)NR₁₀R₁₄, C(NCN)SR₉, (2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-oxadiazolyl[1,2,4]), (2-oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl), (2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl); wherein all of the heterocyclic ring systems may be optionally substituted one or more times by R₁₄;

30 Y' is O or S;

35 R₇ is -(CR₄R₅)_qR₁₂ or C₁₋₆ alkyl wherein the R₁₂ or C₁₋₆ alkyl group is optionally substituted one or more times by C₁₋₂ alkyl optionally substituted by one to three fluorines, -F, -Br, -Cl, -NO₂, -NR₁₀R₁₁, -C(O)R₈, -C(O)OR₈, -OR₈, -CN, -C(O)NR₁₀R₁₁, -OC(O)NR₁₀R₁₁, -OC(O)R₈, -NR₁₀C(O)NR₁₀R₁₁, -NR₁₀C(O)R₁₁, -NR₁₀C(O)OR₉, -NR₁₀C(O)R₁₃, -C(NR₁₀)NR₁₀R₁₁, -C(NCN)NR₁₀R₁₁, -C(NCN)SR₉, -NR₁₀C(NCN)SR₉, -NR₁₀C(NCN)NR₁₀R₁₁, -NR₁₀S(O)₂R₉, -S(O)_mR₉,

-NR₁₀C(O)C(O)NR₁₀R₁₁, -NR₁₀C(O)C(O)R₁₀, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl;

q is 0, 1, or 2;

R₁₂ is C₃₋₇ cycloalkyl, (2-, 3- or 4-pyridyl), pyrimidyl, pyrazolyl, (1- or 2-imidazolyl), thiazolyl, triazolyl, pyrrolyl, piperazinyl, piperidinyl, morpholinyl, furanyl, (2- or 3-thienyl), (4- or 5-thiazolyl), quinolinyl, naphthyl, or phenyl;

R₈ is hydrogen or R₉;

R₈ is R₈ or fluorine;

R₉ is C₁₋₄ alkyl optionally substituted by one to three fluorines;

R₁₀ is OR₈ or R₁₁;

R₁₁ is hydrogen, or C₁₋₄ alkyl optionally substituted by one to three fluorines; or when R₁₀ and R₁₁ are as NR₁₀R₁₁ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom which is O, N, or S;

R₁₃ is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl, imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted or substituted by one or two C₁₋₂ alkyl groups;

R₁₄ is hydrogen or R₇; or when R₁₀ and R₁₄ are as NR₁₀R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom which is O, N, or S;

provided that:

f) when R₁₂ is N-pyrazolyl, N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, N-piperidinyl, or N-morpholinyl, then q is not 1; or

or the pharmaceutically acceptable salts thereof.

Detailed Description of the Invention

This invention relates to the novel compounds of Formula (I), and to pharmaceutical compositions comprising a compound of Formula (I) and a pharmaceutically acceptable carrier or diluent. This invention also relates to a method of mediating or inhibiting the enzymatic activity (or catalytic activity) of PDE IV in a mammal in need thereof and to inhibiting the production of TNF in a mammal in need thereof, which comprises administering to said mammal an effective amount of a compound of Formula (I).

Phosphodiesterase IV inhibitors are useful in the treatment of a variety of allergic and inflammatory diseases including: asthma, chronic bronchitis, atopic dermatitis, urticaria, allergic rhinitis, allergic conjunctivitis, vernal conjunctivitis, eosinophilic granuloma, psoriasis, rheumatoid arthritis, septic shock, ulcerative colitis, Crohn's disease, reperfusion injury of the myocardium and brain, chronic glomerulonephritis, endotoxic shock and adult respiratory distress syndrome. In addition, PDE IV inhibitors are useful in the treatment of diabetes insipidus, [Kidney Int., 37:362, 1990; Kidney Int., 35:494, 1989] and central

nervous system disorders such as depression and multi-infarct dementia.

The compounds of Formula (I) are also useful in the treatment of viral infections, where such viruses are sensitive to upregulation by TNF or will elicit TNF production *in vivo*. The viruses contemplated for treatment herein are those that produce TNF as a result of infection, or those which are sensitive to inhibition, such as by decreased replication, directly or indirectly, by the TNF inhibitors of Formula (1). Such viruses include, but are not limited to HIV-1, HIV-2 and HIV-3, cytomegalovirus (CMV), influenza, adenovirus and the Herpes group of viruses, such as, but not limited to, *Herpes zoster* and *Herpes simplex*.

10 This invention more specifically relates to a method of treating a mammal, afflicted with a human immunodeficiency virus (HIV), which comprises administering to such mammal an effective TNF inhibiting amount of a compound of Formula (I).

The compounds of Formula (I) may also be used in association with the veterinary treatment of animals, other than in humans, in need of inhibition of TNF production. TNF mediated diseases for treatment, therapeutically or prophylactically, in animals include disease states such as those noted above, but in particular viral infections. Examples of such viruses include, but are not limited to feline immunodeficiency virus (FIV) or other retroviral infection such as equine infectious anemia virus, caprine arthritis virus, visna virus, maedi virus and other lentiviruses.

20 The compounds of Formula (I) are also useful in the treatment of yeast and fungal infections, where such yeast and fungi are sensitive to upregulation by TNF or will elicit TNF production *in vivo*. A preferred disease state for treatment is fungal meningitis. Additionally, the compounds of Formula (I) may be administered in conjunction with other drugs of choice for systemic yeast and fungal infections. Drugs of choice for fungal infections, include but are not limited to the class of compounds called the polymyxins, such as Polymycin B, the class of compounds called the imidazoles, such as clotrimazole, econazole, miconazole, and ketoconazole; the class of compounds called the triazoles, such as fluconazole, and itranazole, and the class of compound called the Amphotericins, in particular Amphotericin B and liposomal Amphotericin B.

30 The co-administration of the anti-fungal agent with a compound of Formula (I) may be in any preferred composition for that compound such as is well known to those skilled in the art, for instance the various Amphotericin B formulations. Co-administration of an anti-fungal agent with a compound of Formula (I) may mean simultaneous administration or in practice, separate administration of the agents to the mammal but in a consecutive manner. In particular, the compounds of Formula (I) may be co-administered with a formulation of Amphotericin B, notably for systemic fungal infections. The preferred organism for treatment is the *Candida* organism. The compounds of Formula (I) may be co-administered in a similar manner with anti-viral or anti-bacterial agents.

The compounds of Formula (I) may also be used for inhibiting and/or reducing the toxicity of an anti-fungal, anti-bacterial or anti-viral agent by administering an effective amount of a compound of Formula (I) to a mammal in need of such treatment. Preferably, a compound of Formula (I) is administered for inhibiting or reducing the toxicity of the

5 Amphotericin class of compounds, in particular Amphotericin B.

When R₁ for the compounds of Formula (I) is an alkyl substituted by 1 or more halogens, the halogens are preferably fluorine and chlorine, more preferably a C₁₋₄ alkyl substituted by 1 or more fluorines. The preferred halo-substituted alkyl chain length is one or two carbons, and most preferred are the moieties -CF₃, -CH₂F, -CHF₂, -CF₂CHF₂,
 10 -CH₂CF₃, and -CH₂CHF₂. Preferred R₁ substituents for the compounds of Formula (I) are CH₂-cyclopropyl, CH₂-C₅₋₆ cycloalkyl, C₄₋₆ cycloalkyl, C₇₋₁₁ polycycloalkyl, (3- or 4-cyclopentenyl), phenyl, tetrahydrofuran-3-yl, benzyl or C₁₋₂ alkyl optionally substituted by 1 or more fluorines, -(CH₂)₁₋₃C(O)O(CH₂)₀₋₂CH₃, -(CH₂)₁₋₃O(CH₂)₀₋₂CH₃, and -(CH₂)₂₋₄OH.

15 When the R₁ term contains the moiety (CR₄R₅), the R₄ and R₅ terms are independently hydrogen or alkyl. This allows for branching of the individual methylene units as (CR₄R₅)_n or (CR₄R₅)_m; each repeating methylene unit is independent of the other, e.g., (CR₄R₅)_n wherein n is 2 can be -CH₂CH(-CH₃)-, for instance. The individual hydrogen atoms of the repeating methylene unit or the branching hydrocarbon can optionally be
 20 substituted by fluorine independent of each other to yield, for instance, the preferred R₁ substitutions, as noted above.

When R₁ is a C₇₋₁₁ polycycloalkyl, examples are bicyclo[2.2.1]-heptyl, bicyclo[2.2.2]octyl, bicyclo[3.2.1]octyl, tricyclo[5.2.1.0^{2,6}]decyl, etc. additional examples of which are described in Saccamano *et al.*, WO 87/06576, published 5 November 1987,
 25 which disclosure is incorporated herein by reference in its entirety.

Z is preferably C(O)R₈, C(O)OR₈, C(O)NR₈R₈, C(NR₈)NR₈R₈, CN, C(NOR₈)R₈, C(O)NR₈NR₈C(O)R₈, C(NCN)NR₈R₈, C(NCN)SR₉, (1-, 4- or 5-{R₈}-2-imidazolyl), (1-, 4- or 5-{R₈}-3-pyrazolyl), (1-, 2- or 5-{R₈}-4-triazolyl[1,2,3]), (1-, 2-, 4- or 5-{R₈}-3-triazolyl[1,2,4]), (1- or 2-{R₈}-5-tetrazolyl), (4- or
 30 5-{R₈}-2-oxazolyl), (3- or 4-{R₈}-5-isoxazolyl), (3-{R₈}-5-oxadiazolyl[1,2,4]), (5-{R₈}-3-oxadiazolyl[1,2,4]), (5-{R₈}-2-oxadiazolyl[1,3,4]), (5-{R₈}-2-thiadiazolyl[1,3,4]), (4- or 5-{R₈}-2-thiazolyl), (4- or 5-{R₈}-2-oxazolidinyl), (4- or 5-{R₈}-2-thiazolidinyl), (1-, 4- or 5-{R₈}-2-imidazolidinyl); most preferred are those compounds wherein the R₈ group of Z is R₄.

35 X₅ is preferably hydrogen, C₁₋₂ alkyl optionally substituted by one to three fluorines, OR₈, CN, C(O)R₈, C(O)OR₈, C(O)NR₈R₈, or NR₈R₈.

Preferred X groups for Formula (I) are those wherein X is YR₂ and Y is oxygen. The preferred X₂ group for Formula (I) is that wherein X₂ is oxygen. The preferred X₃

group for Formula (I) is that wherein X₃ is hydrogen. Preferred R₂ groups, where applicable, are C₁₋₂ alkyl optionally substituted by 1 or more halogens. The halogen atoms are preferably fluorine and chlorine, more preferably fluorine. More preferred R₂ groups are those wherein R₂ is methyl, or the fluoro-substituted alkyls, specifically a C₁₋₂ alkyl, such as a -CF₃, -CHF₂, or -CH₂CHF₂ moiety. Most preferred are the -CHF₂ and -CH₃ moieties.

Preferred R₃ moieties are C(O)NH₂, C≡CR₈, CN, C(Z')H, CH₂OH, CH₂F, CF₂H, and CF₃. More preferred are C≡CH and CN. Z' is preferably O or NOR₈.

Preferred R₇ moieties include optionally substituted -(CH₂)₁₋₂(cyclopropyl), -(CH₂)₀₋₂(cyclobutyl), -(CH₂)₀₋₂(cyclopentyl), -(CH₂)₀₋₂(cyclohexyl), -(CH₂)₀₋₂(2-, 3- or 4-pyridyl), -(CH₂)₁₋₂(2-imidazolyl), -(CH₂)₂(4-morpholinyl), -(CH₂)₂(4-piperazinyl), -(CH₂)₁₋₂(2-thienyl), -(CH₂)₁₋₂(4-thiazolyl), and -(CH₂)₀₋₂phenyl;

Preferred rings when R₁₀ and R₁₁ in the moiety -NR₁₀R₁₁ together with the nitrogen to which they are attached form a 5 to 7 membered ring optionally containing at least one additional heteroatom which is O, N, or S include, but are not limited to 1-imidazolyl, 2-(R₈)-1-imidazolyl, 1-pyrazolyl, 3-(R₈)-1-pyrazolyl, 1-triazolyl, 2-triazolyl, 5-(R₈)-1-triazolyl, 5-(R₈)-2-triazolyl, 5-(R₈)-1-tetrazolyl, 5-(R₈)-2-tetrazolyl, 1-tetrazolyl, 2-tetrazolyl, morpholinyl, piperazinyl, 4-(R₈)-1-piperazinyl, or pyrrolyl ring.

Preferred rings when R₁₀ and R₁₄ in the moiety -NR₁₀R₁₄ together with the nitrogen to which they are attached may form a 5 to 7 membered ring optionally containing at least one additional heteroatom which is O, N, or S include, but are not limited to 1-imidazolyl, 1-pyrazolyl, 1-triazolyl, 2-triazolyl, 1-tetrazolyl, 2-tetrazolyl, morpholinyl, piperazinyl, and pyrrolyl. The respective rings may be additionally substituted, where applicable, on an available nitrogen or carbon by the moiety R₇ as described herein for Formula (I). Illustrations of such carbon substitutions includes, but are not limited to, 2-(R₇)-1-imidazolyl, 4-(R₇)-1-imidazolyl, 5-(R₇)-1-imidazolyl, 3-(R₇)-1-pyrazolyl, 4-(R₇)-1-pyrazolyl, 5-(R₇)-1-pyrazolyl, 4-(R₇)-2-triazolyl, 5-(R₇)-2-triazolyl, 4-(R₇)-1-triazolyl, 5-(R₇)-1-triazolyl, 5-(R₇)-1-tetrazolyl, and 5-(R₇)-2-tetrazolyl. Applicable nitrogen substitution by R₇ includes, but is not limited to, 1-(R₇)-2-tetrazolyl, 2-(R₇)-1-tetrazolyl, 4-(R₇)-1-piperazinyl. Where applicable, the ring may be substituted one or more times by R₇.

Preferred groups for NR₁₀R₁₄ which contain a heterocyclic ring are 5-(R₁₄)-1-tetrazolyl, 2-(R₁₄)-1-imidazolyl, 5-(R₁₄)-2-tetrazolyl, or 4-(R₁₄)-1-piperazinyl.

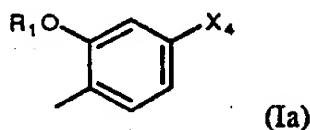
Preferred rings for R₁₃ include (2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-oxadiazolyl[1,2,4]), (2-oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl), (2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl).

When the R₇ group is optionally substituted by a heterocyclic ring such as imidazolyl, pyrazolyl, triazolyl, tetrazolyl, or thiazolyl, the heterocyclic ring itself may be optionally substituted by R₈ either on an available nitrogen or carbon atom, such as 1-(R₈)-2-imidazolyl, 1-(R₈)-4-imidazolyl, 1-(R₈)-5-imidazolyl, 1-(R₈)-3-pyrazolyl, 1-(R₈)-4-pyrazolyl, 1-(R₈)-5-pyrazolyl, 1-(R₈)-4-triazolyl, or 1-(R₈)-5-triazolyl. Where applicable, the ring may be substituted one or more times by R₈.

Preferred are those compounds of Formula (I) wherein R₁ is -CH₂-cyclopropyl, -CH₂-C₅₋₆ cycloalkyl, -C₄₋₆ cycloalkyl, tetrahydrofuran-3-yl, (3- or 4-cyclopentenyl), benzyl or -C₁₋₂ alkyl optionally substituted by 1 or more fluorines, and -(CH₂)₂₋₄ OH; R₂ is methyl or fluoro-substituted alkyl, R₃ is CN or C≡CR₈; and X is YR₂.

Most preferred are those compounds wherein R₁ is -CH₂-cyclopropyl, cyclopentyl, methyl or CF₂H; R₃ is CN or C≡CH; X is YR₂; Y is oxygen; X₂ is oxygen; X₃ is hydrogen; and R₂ is CF₂H or methyl. In most cases the cis configuration is preferred over the trans configuration. However, substituent pattern may influence the overall activity of these compounds in some manner which results in the trans configuration being the more active configuration. While the cis configuration is thought to be more active in most instances, this must be confirmed on for each compound.

A preferred subgenus of the compounds of Formula (I) is the group of compounds of Formula (Ia)

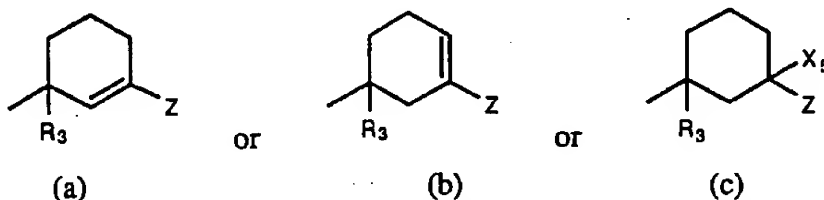


wherein:

R₁ is CH₂-cyclopropyl, CH₂-C₅₋₆ cycloalkyl, C₄₋₆ cycloalkyl, C₇₋₁₁ polycycloalkyl, (3- or 4-cyclopentenyl), phenyl, tetrahydrofuran-3-yl, benzyl or C₁₋₂ alkyl optionally substituted by 1 or more fluorines, -(CH₂)₁₋₃C(O)O(CH₂)₀₋₂CH₃, -(CH₂)₁₋₃O(CH₂)₀₋₂CH₃, and -(CH₂)₂₋₄OH;

X is YR₂, halogen, nitro, NR₄R₅, or formyl amine;

X₄ is



X₅ is H, R₉, OR₈, CN, C(O)R₈, C(O)OR₈, C(O)NR₈R₈, or NR₈R₈;

Y is O or S(O)_m;

m' is 0, 1, or 2;

R₂ is -CH₃ or -CH₂CH₃ optionally substituted by 1 or more halogens;

R₃ is hydrogen, C₁₋₄ alkyl, CH₂NHC(O)C(O)NH₂, halo-substituted C₁₋₄ alkyl, CN, CH₂OR₈, C(Z')H, C(O)OR₈, C(O)NR₈R₁₀, or C≡CR₈;

Z' is O or NOR₈;

5 Z is C(O)R₁₄, C(O)OR₁₄, C(O)NR₁₀R₁₄, C(NR₁₀)NR₁₀R₁₄, CN, C(NOR₈)R₁₄, C(O)NR₈NR₈C(O)R₈, C(O)NR₈NR₁₀R₁₄, C(NOR₁₄)R₈, C(NR₈)NR₁₀R₁₄, C(NR₁₄)NR₈R₈, C(NCN)NR₁₀R₁₄, C(NCN)SR₉, (1-, 4- or 5-{R₁₄}-2-imidazolyl), (1-, 4- or 5-{R₁₄}-3-pyrazolyl), (1-, 2- or 5-{R₁₄}-4-triazolyl[1,2,3]), (1-, 2-, 4- or 5-{R₁₄}-3-triazolyl[1,2,4]), (1- or 2-{R₁₄}-5-tetrazolyl), (4- or 5-{R₁₄}-2-oxazolyl), (3- or 4-{R₁₄}-5-isoxazolyl), (3-{R₁₄}-5-oxadiazolyl[1,2,4]), (5-{R₁₄}-3-oxadiazolyl[1,2,4]), (5-{R₁₄}-2-oxadiazolyl[1,3,4]), (5-{R₁₄}-2-thiadiazolyl[1,3,4]), (4- or 5-{R₁₄}-2-thiazolyl), (4- or 5-{R₁₄}-2-oxazolidinyl), (4- or 5-{R₁₄}-2-thiazolidinyl), (1-, 4- or 5-{R₁₄}-2-imidazolidinyl);

15 R₇ is -(CR₄R₅)_qR₁₂ or C₁₋₆ alkyl wherein the R₁₂ or C₁₋₆ alkyl group is optionally substituted one or more times by C₁₋₂ alkyl optionally substituted by one to three fluorines, -F, -Br, -Cl, -NO₂, -NR₁₀R₁₁, -C(O)R₈, -C(O)OR₈, -OR₈, -CN, -C(O)NR₁₀R₁₁, -OC(O)NR₁₀R₁₁, -OC(O)R₈, -NR₁₀C(O)NR₁₀R₁₁, -NR₁₀C(O)R₁₁, -NR₁₀C(O)OR₉, -NR₁₀C(O)R₁₃, -C(NR₁₀)NR₁₀R₁₁, -C(NCN)NR₁₀R₁₁, -C(NCN)SR₉, -NR₁₀C(NCN)SR₉, -NR₁₀C(NCN)NR₁₀R₁₁, -NR₁₀S(O)₂R₉, -S(O)_mR₉, -NR₁₀C(O)C(O)NR₁₀R₁₁, -NR₁₀C(O)C(O)R₁₀, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl;

q is 0, 1, or 2;

R₁₂ is C₃₋₇ cycloalkyl, (2-, 3- or 4-pyridyl), (1- or 2-imidazolyl), piperazinyl, morpholinyl, (2- or 3-thienyl), (4- or 5-thiazolyl), or phenyl;

25 R₈ is independently selected from hydrogen or R₉;

R₉ is C₁₋₄ alkyl optionally substituted by one to three fluorines;

R₁₀ is OR₈ or R₁₁;

30 R₁₁ is hydrogen or C₁₋₄ alkyl optionally substituted by one to three fluorines; or when R₁₀ and R₁₁ are as NR₁₀R₁₁ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom which is O, N, or S;

R₁₃ is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl, imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted or substituted by one or two C₁₋₂ alkyl groups;

35 R₁₄ is hydrogen or R₇; or when R₁₀ and R₁₄ are as NR₁₀R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing one or more additional heteroatoms which is O, N, or S;

provided that:

a) when R₁₂ is N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, or N-morpholinyl, then q is not 1; or
or the pharmaceutically acceptable salts thereof.

Exemplified compounds of Formula (I) are:

- 5 methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate;, SB 212179
ethyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate;, SB
methyl 5-cyano-5-(3-cyclopentylloxy-4-methoxyphenyl)cyclohex-1-ene-1-
10 carboxylate;, SB 211572
methyl 5-(3,4-bisdifluoromethoxyphenyl)-5-cyanocyclohex-1-ene-1-carboxylate;, SB
methyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate];, SB 210667
15 ethyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate];, SB 211600
methyl *cis*-[3-cyano-3-(3-cyclopentylloxy-4-methoxyphenyl)cyclohexane-1-carboxylate];, SB 212066
methyl *cis*-[3-(3,4-bisdifluoromethoxyphenyl)-3-cyanocyclohexane-1-
20 carboxylate];, SB
cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid];, SB 210984
cis-[3-cyano-3-(3-cyclopentylloxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid]; SB 212510
25 *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide]; SB 211529
cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-N-methylcarboxamide]; SB 213021
cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-N,N-dimethylcarboxamide]; SB 212697
30 *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-{N-(4-bromobenzyl)carboxamide}]; SB 212698
cis-[3-cyano-3-(3-cyclopentylloxy-4-methoxyphenyl)cyclohexane-1-carboxamide]; SB 214236
35 *cis*-[3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1,3-dicyano-cyclohexane]; SB 212188
cis-[3-(3-cyclopentylloxy-4-methoxyphenyl)-1,3-dicyanocyclohexane]; SB 213832

cis-{3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1-(3-methyl[1,2,4]oxadiazol-5-yl)cyclohexane}; SB 213826

cis-{3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)-1-(3-methyl[1,2,4]-oxadiazol-5-yl)cyclohexane}; SB 214243

5 *trans*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate]; SB 213677

trans-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)-cyclohexane-1-carboxylate]; SB 213951

10 *trans*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid]; SB 213731

trans-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid];, SB 213921

trans-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide]; SB 213835

15 *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxamide]; SB 213921

trans-[3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1,3-dicyanocyclohexane]; SB 213920

20 *trans*-[3-(3-cyclopentyloxy-4-methoxyphenyl)-1,3-dicyanocyclohexane]; and SB 214241,

trans-{3-cyano-3-(3-cyclopentyloxyphenyl)-1-(3-methyl[1,2,4]oxadiazol-5-yl)cyclo-hexane};, SB 214242

methyl 3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate;

25 methyl 5-cyano-5-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate;

3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1,3-dicarbonitrile; and

5-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1,5-dicarbonitrile.

30 Some compounds of Formula (I) may exist in both racemic and optically active forms; some may also exist in distinct diastereomeric forms possessing distinct physical and biological properties. All of these compounds are considered to be within the scope of the present invention. Therefore another aspect of the present invention is the administration of either a racemate, a single enantiomeric form, a single diastereomeric form, or mixtures thereof.

35 The terms *cis* and *trans* denote stereochemistry at the C-1 position of the cyclohexane ring relative to the R₃ group at the C-3 position.

The terms "C₁₋₃ alkyl", "C₁₋₄ alkyl", "C₁₋₆ alkyl" or "alkyl" include both straight or branched chain radicals of 1 to 10, unless the chain length is limited thereto, including, but

not limited to methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, isobutyl, *tert*-butyl, and the like. "Alkenyl" includes both straight or branched chain radicals of 1 to 6 carbon lengths, unless the chain length is limited thereto, including but not limited to vinyl, 1-propenyl, 2-propenyl, 2-propynyl, or 3-methyl-2-propenyl. "Cycloalkyl" or "cycloalkyl alkyl" includes groups of 3-7 carbon atoms, such as cyclopropyl, cyclopropylmethyl, cyclopentyl, or cyclohexyl. "Aryl" or "aralkyl", unless specified otherwise, means an aromatic ring or ring system of 6 to 10 carbon atoms, such as phenyl, benzyl, phenethyl, or naphthyl. Preferably the aryl is monocyclic, i.e., phenyl. The alkyl chain includes both straight or branched chain radicals of 1 to 4 carbon atoms. "Heteroaryl" as used herein, is meant an aromatic ring system containing one or more heteroatoms, such as imidazolyl, triazolyl, oxazolyl, pyridyl, pyrimidyl, pyrazolyl, pyrrolyl, furanyl, or thienyl. "Halo" as used herein is meant all halogens, i.e., chloro, fluoro, bromo, or iodo.

The phrase "inhibiting the production of IL-1" or "inhibiting the production of TNF" means:

- a) a decrease of excessive *in vivo* IL-1 or TNF levels, respectively, in a human to normal levels or below normal levels by inhibition of the *in vivo* release of IL-1 by all cells, including but not limited to monocytes or macrophages;
- b) a down regulation, at the translational or transcriptional level, of excessive *in vivo* IL-1 or TNF levels, respectively, in a human to normal levels or below normal levels; or
- c) a down regulation, by inhibition of the direct synthesis of IL-1 or TNF levels as a posttranslational event.

"TNF mediated disease or disease states" means any and all disease states in which TNF plays a role, either by production of TNF itself, or by TNF causing another cytokine to be released, such as but not limited to IL-1 or IL-6. A disease state in which IL-1, for instance is a major component, and whose production or action, is exacerbated or secreted in response to TNF, would therefore be considered a disease state mediated by TNF. As TNF- β (also known as lymphotoxin) has close structural homology with TNF- α (also known as cachectin), and since each induces similar biologic responses and binds to the same cellular receptor, both TNF- α and TNF- β are inhibited by the compounds of the present invention and thus are herein referred to collectively as "TNF" unless specifically delineated otherwise. Preferably TNF- α is inhibited.

"Cytokine" means any secreted polypeptide that affects the functions of cells, and is a molecule which modulates interactions between cells in immune, inflammatory, or hematopoietic responses. A cytokine includes, but is not limited to, monokines and lymphokines regardless of which cells produce them. For instance, a monokine is generally referred to as being produced and secreted by a mononuclear cell, such as a macrophage and/or monocyte, but many other cells produce monokines, such as natural killer cells, fibroblasts, basophils, neutrophils, endothelial cells, brain astrocytes, bone marrow stromal

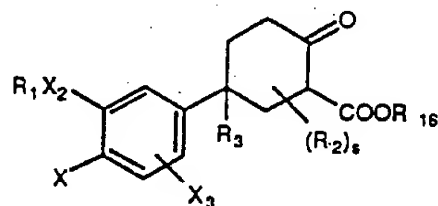
cells, epidermal keratinocytes, and B-lymphocytes. Lymphokines are generally referred to as being produced by lymphocyte cells. Examples of cytokines for the present invention include, but are not limited to, Interleukin-1 (IL-1), Interleukin-6 (IL-6), Interleukin-8 (IL-8), Tumor Necrosis Factor-alpha (TNF- α) and Tumor Necrosis Factor-beta (TNF- β).

The cytokine inhibited by the present invention for use in the treatment of a HIV-infected human must be a cytokine which is implicated in (a) the initiation and/or maintenance of T cell activation and/or activated T cell-mediated HIV gene expression and/or replication, and/or (b) any cytokine-mediated disease associated problem such as cachexia or muscle degeneration. Preferably this cytokine is TNF- α .

All of the compounds of Formula (I) are useful in the method of inhibiting the production of TNF, preferably by macrophages, monocytes or macrophages and monocytes, in a mammal, including humans, in need thereof. All of the compounds of Formula (I) are useful in the method of inhibiting or mediating the enzymatic or catalytic activity of PDE IV and in treatment of disease states mediated thereby.

METHODS OF PREPARATION:

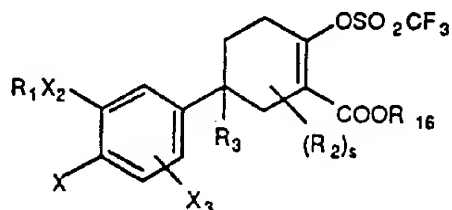
Preparing compounds of Formula (I) can be carried out by one of skill in the art according to the procedures outlined in the Examples, *infra*. For example, reacting a compound of Formula (2)



(2)

wherein X or X_3 is other than Br, I, NO_2 , amino, or $S(O)_mR_2$ when m ' is 0, 1 or 2, R_1 represents R_1 as defined in relation to Formula (I) or a group convertible to R_1 and X , X_2 and X_3 represent X , X_2 and X_3 as defined in relation to Formula (I) or a group convertible to X , X_2 or X_3 and R_3 represents R_3 as defined in relation to Formula (I) or a group convertible to R_3 and R_{16} is an alkyl, phenyl or benzyl group with, e.g.,

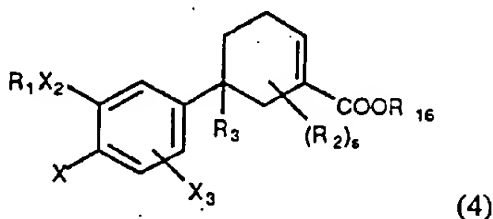
trifluoromethanesulfonic anhydride in the presence of a suitable base, such as a hindered amine base, in a suitable solvent, such as dichloromethane, provides a compound of Formula (3)



(3)

wherein X or X_3 is other than Br, I, NO_2 , amino, or $S(O)_mR_2$ when m ' is 0, 1 or 2, R_1

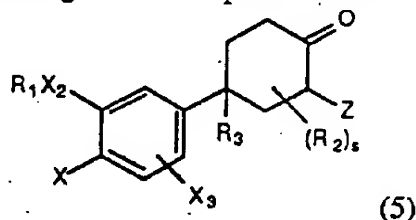
represents R_1 as defined in relation to Formula (I) or a group convertible to R_1 and X , X_2 and X_3 represent X , X_2 and X_3 as defined in relation to Formula (I) or a group convertible to X , X_2 or X_3 and R_3 represents R_3 as defined in relation to Formula (I) or a group convertible to R_3 and R_{16} is an alkyl, phenyl or benzyl group. Palladium-catalyzed
 5 reduction of such a compound of Formula (3) under suitable conditions then provides a compound of Formula (4)



a subset of the compounds of Formula (1) wherein X or X_3 is other than Br , I , NO_2 , amino, or $S(O)_{m'}R_2$ when m' is 0, 1 or 2, the double bond is present, R_1 represents R_1 as defined
 10 in relation to Formula (I) or a group convertible to R_1 and X , X_2 and X_3 represent X , X_2 and X_3 as defined in relation to Formula (I) or a group convertible to X , X_2 or X_3 and R_3 represents R_3 as defined in relation to Formula (I) or a group convertible to R_3 and R_{16} is an alkyl, phenyl or benzyl group. Such compounds of Formula (1) wherein X or X_3 is other
 15 than Br , I , NO_2 , amino, or $S(O)_{m'}R_2$ when m' is 0, 1 or 2, the double bond is present and Z is $COOR_{16}$ can be converted to other compounds of Formula (1) wherein X or X_3 is other than Br , I , NO_2 , amino, or $S(O)_{m'}R_2$ when m' is 0, 1 or 2 and the double bond is present by standard procedures well known in the art [with proper manipulation
 (protection/deprotection) of any chemically sensitive functional groups, if necessary] to the corresponding ester, amide, nitrile, oxazolidinone, etc., Z groups of Formula (I) wherein X
 20 or X_3 is other than Br , I , NO_2 , amino, or $S(O)_{m'}R_2$ when m' is 0, 1 or 2 and the double bond is present. Alternatively, such compounds of Formula (I) wherein X or X_3 is other than Br , I , NO_2 , amino, or $S(O)_{m'}R_2$ when m' is 0, 1 or 2, the double bond is present, R_1 represents R_1 as defined in relation to Formula (I) or a group convertible to R_1 and X , X_2
 25 and X_3 represent X , X_2 and X_3 as defined in relation to Formula (I) or a group convertible to X , X_2 or X_3 and R_3 represents R_3 as defined in relation to Formula (I) or a group convertible to R_3 and R_{16} is an alkyl, phenyl or benzyl group may then be reduced by, e.g., catalytic hydrogenation, to compounds of Formula (I) wherein X or X_3 is other than Br , I , NO_2 , amino, or $S(O)_{m'}R_2$ when m' is 0, 1 or 2 and the double bond is absent. Functional
 30 conversion of the Z group in such compounds of Formula (I) wherein X or X_3 is other than Br , I , NO_2 , amino, or $S(O)_{m'}R_2$ when m' is 0, 1 or 2, the double bond is absent, R_1 represents R_1 as defined in relation to Formula (I) or a group convertible to R_1 and X , X_2 and X_3 represent X , X_2 and X_3 as defined in relation to Formula (I) or a group convertible
 35 to X , X_2 or X_3 and R_3 represents R_3 as defined in relation to Formula (I) or a group convertible to R_3 and R_{16} is an alkyl, phenyl or benzyl group to other Z groups can be accomplished by standard procedures well known in the art [with proper manipulation

(protection/deprotection) of any chemically sensitive functional groups, if necessary]. For example, preparation of some compounds of Formula (I) wherein R_3 is, e.g., $C(=Z')H$ proceed in an analogous fashion from the compound of Formula (2) wherein $=Z'$ is an aldehyde protecting group, such as a dimethylacetal or a dioxolane, followed by aldehyde deprotection and subsequent manipulation by standard procedures known to those of skill in the art to the remaining compounds of Formula (I). Likewise, isomerization of compounds of Formula (I) wherein X or X_3 is other than Br, I, NO_2 , amino, or $S(O)_mR_2$ when m is 0, 1 or 2, the double bond is absent, R_1 represents R_1 as defined in relation to Formula (I) or a group convertible to R_1 and X ; X_2 and X_3 represent X , X_2 and X_3 as defined in relation to Formula (I) or a group convertible to X , X_2 or X_3 and R_3 represents R_3 as defined in relation to Formula (I) or a group convertible to R_3 and R_{16} is an alkyl, phenyl or benzyl group and the R_3 and Z groups are *cis* to the isomer wherein the R_3 and Z groups are *trans* can be accomplished either under kinetic or thermodynamic deprotonation conditions by standard procedures well known in the art with proper manipulation (protection/deprotection) of any chemically sensitive functional groups.

Alternatively, compounds of Formula (I) wherein X or X_3 is other than Br, I, NO_2 , amino, or $S(O)_mR_2$ when m is 0, 1 or 2 may be prepared by a reaction sequence analogous to that described above but starting with a compound of Formula (5)



wherein X or X_3 is other than Br, I, NO_2 , amino, or $S(O)_m$ when m is 0, 1 or 2, the double bond is absent, R_1 represents R_1 as defined in relation to Formula (I) or a group convertible to R_1 and X , X_2 and X_3 represent X , X_2 and X_3 as defined in relation to Formula (I) or a group convertible to X , X_2 or X_3 and R_3 represents R_3 as defined in relation to Formula (I) or a group convertible to R_3 and Z represents Z as defined in relation to Formula (I) or a group convertible to Z .

Any remaining compounds of Formula (I) not described therein may be prepared by the analogous processes disclosed herein which comprise:

With proper manipulation (protection/deprotection) of any chemically sensitive functional groups:

a) Compounds of Formula (I) wherein X or X_3 are formyl amine may be formed at the last step, by formylating a compound wherein X or X_3 is NH_2 , obtained by removal of a protecting group from the amine functionality; such protective groups are well known to those skilled in the art, See Greene, T. and Wuts, P.G.M., *Protecting Groups in Organic Synthesis*, 2nd Ed., John Wiley and Sons, New York (1991).

b) Compounds of Formula (I) wherein X or X_3 are Br, I or SR_2 may be

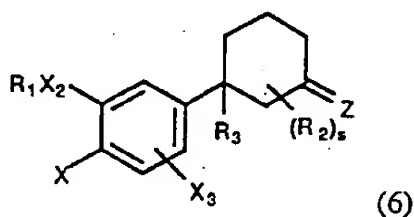
prepared from a similarly deprotected amine by diazotization of the amine and diazonium displacement.

c) Compounds of Formula (I) wherein X or X₃ are NO₂ may be prepared from a similarly deprotected amine by oxidizing the amine to the nitro group.

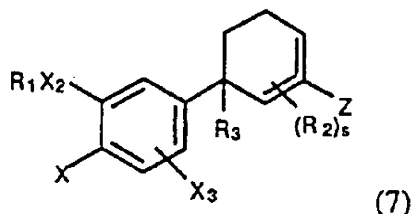
5 d) Compounds of Formula (I) wherein Y is S(O)_{m'} when m' is 1 or 2 may be prepared from the compounds of Formula (I) wherein Y is S by first protecting any other oxidizable groups which are not to be oxidized then oxidizing the SR₂ moiety under conditions well known those skilled in the art and then deprotecting the previously protected group.

10 Compounds of Formulas (2) and (5) may be prepared in turn by the processes described in co-pending applications described in PCT applications PCT/US93/02230, and its predecessor applications PCT/US93/02046, USSN 07/968,806 filed 30 October 1992 and USSN 07/862,114 filed 2 April 1992; and PCT application PCT/US93/10325 filed 12 March 1993 and the predecessor cases from which priority is claimed therein.

15 Alternatively, reacting a compound of the Formula (6)



20 wherein R₁ represents R₁ as defined in relation to Formula (I) or a group convertible to R₁ and X, X₂ and X₃ represent X, X₂ and X₃ as defined in relation to Formula (I) or a group convertible to X, X₂ or X₃ and R₂ and R₃ represent R₂ and R₃ as defined in relation to Formula (I) or a group convertible to R₂ or R₃ and wherein X or X₃ is other than Br, I, NO₂, amino, or S(O)_{m'}R₂ when m' is 0, 1 or 2 and R₃ is other than C(=Z')H, with a suitable base in a suitable non-reacting solvent followed by reaction with a
25 suitable acylating agent, such as LS(O)₂CF₃, wherein L is a leaving group, to provide compounds of the Formula (7)



30 wherein Z is S(O)₂CF₃ and R₃ is other than C(=Z')H. Metal catalyzed carboxymethylation of such compounds of the Formula (7) then provides the compounds of the Formula (I) wherein Z is COOR₁₆, which may be converted to other compounds of the formula (I) as described above. Preparation of such compounds of Formula (1)

wherein R₃ is C(=Z')H proceeds in an analogous fashion from the compound of Formula (6) wherein =Z' is an aldehyde protecting group, such as a dimethylacetal or a dioxolane, followed by deprotection to the aldehyde at the end of the reaction sequence and subsequent elaboration by standard procedures known to those of skill in the art to the remaining compounds of Formula (1) wherein Z' is other than O.

Compounds of the Formula (6) may be prepared in turn by processes described in co-pending application USSN 08/130215 filed 1 October 1993. P50199.

It will be recognized that compounds of Formula (I) may exist in two distinct diastereomeric forms possessing distinct physical and biological properties; such isomers may be separated by standard chromatographic methods.

The following examples and methods are provided to illustrate how to make and use the invention. These materials are not intended to limit the invention in any manner; please refer to the claims appended hereto for determining what has been reserved to the inventors hereunder.

EXAMPLES

Example 1

Preparation of methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate. SB 212179

Route A

1a) Methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)-2-(trifluoromethylsulfonato)-cyclohex-1-ene-1-carboxylate

To a solution of 2,6-di-tert-butyl-4-methylpyridine (10.3 grams [hereinafter g], 50.2 millimoles [hereinafter mmol]) and trifluoromethanesulfonic anhydride (7.07 milliliters [hereinafter mL], 41.8 mmol) in dichloromethane (165 mL) at room temperature under an argon atmosphere was added over 0.5 hours [hereinafter h] a solution of 2-carbomethoxy-4-cyano-4-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexan-1-one (12 g, 33.6 mmol) in dichloromethane (160 mL). The resulting mixture was stirred overnight and then was concentrated to half-volume. Ether was added, the salt was removed by filtration and the filtrate was concentrated *in vacuo*. Purification by flash chromatography, eluting with 3:1 hexanes/ethyl acetate, afforded a yellow oil (14.7 g, 89%). ¹H NMR (400 MHz, CDCl₃) δ 7.0 (dd, J=8.5 and 2 Hz, 1H), 6.95 (d, J=2 Hz, 1H), 6.90 (d, J=8.5 Hz, 1H), 3.85 (m, 8H), 3.10 (AB system, J=15 Hz, 2H), 2.88 (m, 1H), 2.56 (m, 1H), 2.33 (m, 2H), 1.35 (m, 1H), 0.67 (dt, J=7 and 7 Hz, 2H), 0.49 (dt, J=7 and 7 Hz, 2H).

1b) Methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate

To a mixture of methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)-2-(trifluoromethylsulfonato)cyclohex-1-ene-1-carboxylate (0.34 g, 0.69 mmol),

triphenylphosphine (7.2 mg, 0.03 mmol), palladium acetate (3 mg, 0.013 mmol) and tributylamine (0.49 mL, 2.06 mmol) in N,N-dimethylformamide (1.5 mL) was added dropwise formic acid (0.054 mL, 1.37 mmol). The resulting mixture was heated at 60°C under an argon atmosphere for 1 h. The mixture was poured into 2 N hydrochloric acid and was extracted twice with ethyl acetate. The organic extract was washed twice with 2 N hydrochloric acid, once with aqueous sodium bicarbonate, once with brine and was evaporated. Purification by flash chromatography, eluting with 4:1 hexanes/ethyl acetate, provided an oil. Anal. (C₂₀H₂₃NO₄·3/4 H₂O) calcd: C 67.68, H 6.96, N 3.95; found: C 67.50, H 6.78, N 3.80.

Route B:

1c) 3-(3-Cyclopentyloxy-4-methoxyphenyl)cyclohex-2-en-1-one

n-Butyllithium (2.5M in hexanes, 15.5 mL, 38.9 mmol) was added dropwise over 30 min to a solution of 3-cyclopentyloxy-4-methoxybromobenzene (10 g, 37 mmol,) in dry tetrahydrofuran (100 mL) at -78°C under an argon atmosphere. After 1.5 h. this solution was cannulated into a solution of 3-methoxycyclohex-2-enone (4.62g, 37.4 mmol, prepared as in Pearson, A.J.; Richards, I.C.; Gardner, D.V. J. Org. Chem. 1984, 49, 3887-3891) in dry tetrahydrofuran (50 mL) at 0°C under an argon atmosphere. After 2h at room temperature, a mixture of ether and water was added, the aqueous layer was twice more extracted with ether, the combined extract was washed with water and brine, was dried (magnesium sulfate) and was evaporated. Trituration from ether/hexanes provided an off-white solid (7.33 g, 68%). Further purification of the mother liquor by flash chromatography, eluting with 1:3 ethyl acetate/hexanes, followed by trituration from ether/hexanes, provided a white solid (1.59 g, 7%). mp 89-90°C; Anal. (C₁₈H₂₂O₃·1/8 H₂O) calcd: C 74.91, H 7.77; found: C 74.96, H 7.76.

1d) 3-Cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexan-1-one

To a solution of 3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-2-en-1-one (1.46 g, 5.10 mmol) in dry toluene (45 mL) at room temperature under an argon atmosphere was added over 5 min diethylaluminumcyanide (1.0 M solution in toluene, 15.5 mL, 15.5 mmol). After 6h, the reaction was carefully quenched with sodium hydroxide (2 N, 75 mL, 150 mmol), was extracted three times with methylene chloride, the extract was dried (magnesium sulfate) and was evaporated. Purification by flash chromatography, eluting with 1:4 ethyl acetate/hexanes, provided a pale yellow solid (1.20 g, 75%). mp 110-111°C; Anal. (C₁₉H₂₃NO₃·1/8 H₂O) calcd: C 72.30, H 7.42, N 4.44; found: C 72.24, H 7.45, N 4.58.

1e) Methyl 3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate and methyl 5-cyano-5-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate

A mixture of 3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)-cyclohex-1-en-1-yl

trifluoromethanesulfonamide and 5-cyano-5-(3-cyclopentyloxy-4-methoxyphenyl)-cyclohex-1-en-1-yl trifluoromethanesulfonamide (15.9 g, 35.7 mmol), triphenylphosphine (1.87 g, 7.14 mmol), palladium acetate (400 mg, 1.79 mmol) and tributylamine (25.5 mL, 107 mmol) in methanol (300 mL) was saturated with carbon monoxide, then stirred under a carbon monoxide balloon for 4 days. The solvent was evaporated, the residue was diluted with water and was extracted three times with dichloromethane, was dried (magnesium sulfate) and was evaporated. Purification by flash chromatography, eluting with 85:15 hexanes/ethyl acetate, provided a white, waxy solid (6.21 g, 51%, R_f = 0.36 (2:8 ethyl acetate:hexanes)). Also isolated was the isomer: methyl 3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate as a colorless oil (5.01 g, 39%, R_f = 0.43 (2:8 ethyl acetate:hexanes)).

1f) Methyl cis-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylate]

To a solution of methyl 5-cyano-5-(3-cyclopentyloxy-4-methoxyphenyl)-cyclohex-1-ene-1-carboxylate (6.21 g, 17.5 mmol) in methanol (50 mL) was added 10% palladium on activated carbon (0.60 g) and the resulting mixture was hydrogenated at 50 psi for 5h. The mixture was filtered through a pad of Celite[®], the solid was washed with dichloromethane and the filtrate was concentrated under reduced pressure. Purification by flash chromatography, eluting with 85:15 hexanes:ethyl acetate, provided methyl cis-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylate] as a colorless oil (5.60 g, 90%).

Alternatively, to a solution of methyl 3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate (0.10 g, 0.28 mmol) in methanol (10 mL) was added 10% palladium on activated carbon (0.05 g) and the resulting mixture was hydrogenated at 50 psi for 6h. The mixture was filtered through a pad of Celite[®], the solid was washed with dichloromethane and the filtrate was concentrated under reduced pressure. Purification by flash chromatography, eluting with 9:1 hexanes:ethyl acetate, provided methyl cis-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylate] as a colorless oil (0.04 g, 43%). Also isolated was unreacted starting material (0.04g, 42%).

Example 2

Preparation of ethyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate, SB

Following the procedure of Example 1(a)-1(b), except substituting 2-carboethoxy-4-cyano-4-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexan-1-one for 2-carbomethoxy-4-cyano-4-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexan-1-one, the title compound was prepared as an oil (0.57 g, 95%).

Example 3Preparation of methyl 5-cyano-5-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate. SB 211572

Following the procedure of Example 1(a)-1(b), except substituting 2-carbomethoxy-4-cyano-4-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexan-1-one for 2-carbomethoxy-4-cyano-4-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexan-1-one, the title compound was prepared as a white solid (0.23 g, 54%). mp 68-71°C; Anal. (C₂₁H₂₅NO₄·1/2 H₂O) calcd: C 69.40 H 6.93, N 3.85; found: C 69.48, H 7.33, N 3.69.

Example 4

10 Preparation of methyl -5-(3,4-bisdifluoromethoxyphenyl)-5-cyanocyclohex-1-ene-1-carboxylate. SB

Following the procedure of Example 1(a)-1(b), except substituting 2-carbomethoxy-4-(3,4-bisdifluoromethoxyphenyl)-4-cyanocyclohexan-1-one for 2-carbomethoxy-4-cyano-4-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexan-1-one, the title compound was prepared (0.10g, 51%). ¹H NMR (400 MHz, CDCl₃) δ 7.41 (dd, J=8.5 and 2 Hz, 1H), 7.38 (d, J=2 Hz, 1H), 7.32 (d, J=8.5 Hz, 1H), 7.17 (m, 1H), 6.56 (t, J = 73 Hz, 1H), 6.55 (t, J = 73 Hz, 1H), 3.78 (s, 3H), 2.86 (AB system, J=18 Hz, 2H), 2.70 (m, 1H), 2.52 (m, 1H,), 2.24 (m, 1H,), 2.04 (m, 1H).

Example 5

20 Preparation of methyl cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate]. SB 210667

To a solution of methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohex-1-ene-1-carboxylate (0.4 g, 1.17 mmol) in methanol (10 mL) was added 10% palladium on activated carbon (0.15 g) and the resulting mixture was hydrogenated at 50 psi for 3 h. The mixture was filtered through a pad of Celite®, the solid was washed with dichloromethane and the filtrate was concentrated under reduced pressure. Purification by flash chromatography, eluting with 3:1 hexanes/ethyl acetate, provided a colorless oil (0.27 g, 67%). Anal. (C₂₀H₂₅NO₄) calcd: C 69.95, H 7.34, N 4.08; found: C 69.74, H 7.17, N 4.06.

Example 6

30 Preparation of ethyl cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate]. SB 211600

Following the procedure of Example 5, except substituting ethyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate for methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate, the title compound was prepared (0.43 g, 75%). Anal. (C₂₁H₂₇NO₄·1/8 H₂O) calcd: C 70.12, H 7.64, N 3.89; found: C 70.11, H 7.58, N 4.23.

Example 7

Preparation of methyl *cis*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylate], SB 212066

Following the procedure of Example 5, except substituting methyl 5-cyano-5-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate for methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate, the title compound was prepared as a colorless oil (0.1 g, 67%). Anal. (C₂₁H₂₇NO₄·1/2 H₂O) calcd: C 68.83, H 7.70, N 3.82; found: C 68.83, H 7.49, N 3.63.

Example 8

10 Preparation of methyl *cis*-[3-(3,4-bisdifluoromethoxyphenyl)-3-cyanocyclohexane-1-carboxylate]. SB

Following the procedure of Example 5, except substituting methyl 5-cyano-5-(3,4-bisdifluoromethoxyphenyl)cyclohex-1-ene-1-carboxylate for methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate, the title compound was prepared.

Example 9

Preparation of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid] and separation of the 1R,3S- and 1S,3R-enantiomers. SB 210984

20 To a solution of methyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylate] (0.2 g, 0.58 mmol) in methanol (5.8 mL) under an argon atmosphere was added a solution of potassium hydroxide (0.098 g, 1.5 mmol) in water (3.4 mL). The resulting mixture was stirred at room temperature overnight, then poured into acidic water and extracted three times with ethyl acetate. The
25 extract was dried (magnesium sulfate) and concentrated under reduced pressure. Purification by flash chromatography, eluting with 5% methanol/chloroform, provided a foam (0.16 g, 84%). Anal. (C₁₉H₂₃NO₄·1/8H₂O) calcd: C 68.81, H 7.07, N 4.22; found: C 68.81, H 7.17, N 4.26. Chiral separation of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid] was accomplished using preparative
30 HPLC conditions with a 21.2 x 250 mm Chiralpak AD column. The mobile phase of 85:15:0.2 hexanes:isopropanol/water eluted at a flow rate of 10 mL/min with injection of 0.1 g/10 mL at ambient temperature. Ultraviolet detection of the eluting product was employed at 294 nm. Retention times were 26.3 min for the 1R,3S-isomer and 34.2 min for the 1S,3R-isomer.

35 Example 10

Preparation of *cis*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid]. SB 212510

Following the procedure of Example 9, except substituting methyl *cis*-[3-cyano-3-

(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylate] for methyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylate] and using a solvent mixture of 5:5:2 tetrahydrofuran/methanol/water, the title compound was prepared as a white solid (0.02g, 78%). mp 48-50°C; Anal. (C₂₀H₂₅NO₄·5/4 H₂O) calcd: C 66.05, H 7.55, N 3.85; found: C 65.98, H 7.71, N 4.21.

Example 11

Preparation of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide] SB 211529

A solution of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylic acid] (0.1 g, 0.3 mmol) and N-methylmorpholine (0.04 mL, 0.36 mmol) in 1,2-dimethoxyethane (3.2 mL) at room temperature under an argon atmosphere was treated with isobutyl chloroformate (0.045 mL, 0.35 mmol). After 15 min, concentrated ammonium hydroxide (6 drops) was added and the mixture was stirred for an additional 0.5 h. The mixture was partitioned between dichloromethane and 5% aqueous sodium bicarbonate, was extracted three times, the organic extract was dried (potassium carbonate) and the solvent was evaporated. Purification by flash chromatography, eluting with 5% methanol/chloroform, provided a foam (0.05 g, 51%). This material was combined with the product of a reaction conducted on a similar scale and was re-chromatographed. Anal. (C₁₉H₂₄N₂O₃·1/2H₂O) calcd: C 67.63, H 7.47, N 8.30; found: C 67.59, H 7.28, N 8.04.

Example 12

Preparation of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-N-methylcarboxamide] SB 213021

Following the procedure of Example 11, except substituting methylamine hydrochloride for ammonium hydroxide and using a four-fold excess of N-methylmorpholine, the title compound was prepared as a foam. Anal. (C₂₀H₂₆N₂O₃·1/2H₂O) calcd: C 68.35, H 7.74, N 7.97; found: C 68.27, H 7.70, N 7.91.

Example 13

Preparation of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-N,N-dimethylcarboxamide] SB 212697

Following the procedure of Example 11, except substituting dimethylamine hydrochloride for ammonium hydroxide, the title compound was prepared as a white solid. mp 105°C; Anal. (C₂₁H₂₈N₂O₃·1/4H₂O) calcd: C 69.88, H 7.96 N 7.76; found: C 69.72, H 7.78, N 7.74.

Example 14

Preparation of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-(N-(4-bromobenzyl)carboxamide)] SB 212698

Following the procedure of Example 11, except substituting 4-bromobenzyl

amine hydrochloride for ammonium hydroxide and using a four-fold excess of N-methylmorpholine, the title compound was prepared as a white solid. mp 162-163°C; Anal. (C₂₆H₂₉BrN₂O₃·1/2H₂O) calcd: C 61.66, H 5.97 N 5.53; found: C 61.66, H 5.89, N 5.57.

5

Example 15

Preparation of cis-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxamide] SB 214236

Following the procedure of Example 11, except substituting *cis*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid] for *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid], the title compound was prepared as a white solid (0.6g, 98%). mp 145-146°C; Anal. (C₂₀H₂₆N₂O₃·1/8H₂O) calcd: C 69.69, H 7.68 N 8.13; found: C 69.53, H 7.64, N 8.03.

Example 16

15 Preparation of cis-[3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1,3-dicyano-cyclohexane] SB 212188

A solution of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxamide] (0.29 g, 0.87 mmol) and pyridine (0.15 mL, 1.86 mmol) in tetrahydrofuran (5 mL) at room temperature under an argon atmosphere was treated with trifluoroacetic anhydride (0.13 mL, 0.92 mmol). After stirring for 2.5 h, saturated aqueous ammonium chloride was added and the mixture was partitioned between ethyl acetate and 10% hydrochloric acid. The aqueous layer was extracted three times, the organic extract was dried (magnesium sulfate) and evaporated. Purification by flash chromatography, eluting with 2:1 hexanes/ethyl acetate, provided a solid (0.19 g, 71%). mp 138.5-139°C; Anal. (C₁₉H₂₂N₂O₂) calcd: C 73.52, H 7.14, N 9.03; found: C 73.63, H 7.15, N 8.84.

Example 17

Preparation of cis-[3-(3-cyclopentyloxy-4-methoxyphenyl)-1,3-dicyanocyclohexane] SB 213832

30 Following the procedure of Example 16, except substituting *cis*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxamide] for *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide], the title compound was prepared as a white solid (0.18 g, 71%). mp 113-115°C; Anal. (C₂₀H₂₄N₂O₂) calcd: C 74.05, H 7.46, N 8.63; found: C 73.96, H 7.42, N 8.64.

35

Example 18

Preparation of cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1-(3-methyl[1,2,4]oxadiazol-5-yl)cyclohexane] SB 213826

A solution of *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-

methoxyphenyl)cyclohexane-1-carboxamide] (0.097 g, 0.30 mmol) in N,N-dimethylacetamide dimethyl acetal (1.2 mL) was heated at 110°C under an argon atmosphere for 1 h, was cooled and the solvent was evaporated. Dioxane (1.2 mL), acetic acid (1.2 mL), hydroxylamine hydrochloride (0.03 g, 0.42 mmol) and 10% aqueous sodium hydroxide (0.18 mL, 0.45 mmol) were added and the mixture was heated at 90°C under an argon atmosphere for 2 h. The mixture was cooled, water was added, the mixture was extracted three times with methylene chloride, the organic extract was dried (magnesium sulfate) and was evaporated. Purification by flash chromatography, eluting with 35% ethyl acetate/hexanes, provided an oil (0.06 g, 55%). Anal. (C₂₁H₂₅N₃O₃ · 1/4 H₂O) calcd: C 67.81, H 6.91 N 11.30; found: C 67.73, H 6.98, N 11.14.

Example 19

Preparation of *cis*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)-1-(3-methyl[1,2,4]-oxadiazol-5-yl)cyclohexane] SB 214243

Following the procedure of Example 18, except substituting *cis*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxamide] for *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide], the title compound was prepared as a colorless oil (0.29 g, 66%). Anal. (C₂₂H₂₇N₃O₃ · 1/2 H₂O) calcd: C 67.67, H 7.23, N 10.76; found: C 67.72, H 6.94, N 10.52.

Example 20

Preparation of methyl *trans*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate] SB 213677

A solution of methyl *cis* [3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate] (0.82 g, 2.39 mmol) in N-methylpyrrolidinone (10 mL) was treated with sodium cyanide (0.12 g, 2.51 mmol) and heated at 90°C under an argon atmosphere for 6 h. The mixture was cooled, water (30 mL) was added, the mixture was extracted once with ether, once with ethyl acetate, the combined extract was washed with water, was dried (magnesium sulfate) and was evaporated. Purification by flash chromatography, eluting with 3:1 hexanes/ethyl acetate, provided an oil (0.41 g, 50%). Anal. (C₂₀H₂₅NO₄ · 1/8 H₂O) calcd: C 69.49, H 7.36, N 4.05; found: C 69.45, H 7.20, N 4.14.

Example 21

Preparation of methyl *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)-cyclohexane-1-carboxylate] SB 213951

Following the procedure of Example 20, except substituting methyl *cis*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylate] for methyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylate], the title compound was prepared (0.19 g, 40%). Anal. (C₂₁H₂₇NO₄ · 3/4 H₂O) calcd: C 67.99, H 7.74, N 3.76; found: C 68.07, H 7.50, N 4.09.

Example 22

Preparation of *trans*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid]. SB 213731

Following the procedure of Example 9, except substituting methyl *trans*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylate] for methyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylate], the title compound was prepared (0.17 g, 90%). Anal. (C₁₉H₂₂NO₄·Na) calcd: C 64.95, H 6.31, N 3.99; found: C 65.01 H 6.51, N 3.85.

Example 23

10 Preparation of *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid]. SB 213921

Following the procedure of Example 9, except substituting methyl *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylate] for methyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylate] and using
15 a solvent mixture of 5:5:2 tetrahydrofuran/methanol/water, the title compound was prepared (0.075 g, 84%). Anal. (C₂₀H₂₅NO₄·1/2 H₂O) calcd: C 68.16, H 7.44, N 3.97; found: C 67.84, H 7.23, N 4.13.

Example 24

20 Preparation of *trans*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide] SB 213835

A solution of *trans*[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid] (0.19 g, 0.58 mmol) and triethylamine (0.10 mL, 0.70 mmol) in dichloromethane (5.0 mL) at 0°C under an argon atmosphere was treated with isobutyl chloroformate (0.083 mL, 0.70 mmol). After
25 10 min., anhydrous ammonia was condensed into the reaction mixture and the mixture was stirred for an additional 0.5 h while warming to room temperature. The mixture was partitioned between 95:5 dichloromethane/methanol and 5% aqueous sodium carbonate, was extracted three times, the organic extract was dried (magnesium sulfate) and the solvent was evaporated. The residue was recrystallized
30 from ethyl acetate/hexanes to provide a white solid (0.14 g, 76%). mp 174-176°C; Anal. (C₁₉H₂₄N₂O₃) calcd: C 69.49, H 7.37, N 8.53; found: C 69.19, H 7.38, N 8.59.

Example 25

35 Preparation of *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxamide] SB 213921

Following the procedure of Example 11, except substituting *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid] for *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid], the title compound

was prepared as a white solid (0.65 g, 95%). mp 150-151°C; Anal. (C₂₀H₂₆N₂O₃) calcd: C 70.15, H 7.65, N 8.18; found: C 69.90, H 7.66, N 8.04.

Example 26

Preparation of *trans*-[3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1,3-dicyano-
5 cyclohexane] SB 213920

Following the procedure of Example 16, except substituting *trans*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide] for *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide], the title compound was prepared (0.10 g, 92%). Anal. (C₁₉H₂₂N₂O₂) calcd: C 73.52, H 7.16, N 9.03; found: C 73.34, H 7.16, N 9.02.

Example 27

Preparation of *trans*-[3-(3-cyclopentyloxy-4-methoxyphenyl)-1,3-
15 dicyanocyclohexane] SB 214241

Following the procedure of Example 16, except substituting *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxamide] for *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide], the title compound was prepared as a white solid (0.16 g, 66%). mp 118-119°C; Anal. (C₂₀H₂₄N₂O₂·1/8 H₂O) calcd: C 73.53, H 7.48, N 8.58; found: C 73.47, H 7.44, N 8.46.

Example 28

20 Preparation of *trans*-[3-cyano-3-(3-cyclopentyloxyphenyl)-1-(3-
methyl[1,2,4]oxadiazol-5-yl)cyclohexane] SB 214242

Following the procedure of Example 18, except substituting *trans*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide] for *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide], the title compound was prepared as a colorless oil (0.21 g, 70%). Anal. (C₂₂H₂₇N₃O₃·3/2 H₂O) calcd: C 64.69, H 7.03, N 10.29; found: C 64.98, H 6.76, N 9.90.

Example 29

Preparation of 5-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1,5-
30 dicarbonitrile

A suspension of 3-cyano-3-(cyclopentyloxy-4-methoxyphenyl)cyclohexan-1-one (0.63 g, 2.0 mmol), 18-crown-6 (0.006 g, 0.1 mmol) and potassium cyanide (0.006 g, 0.1 mmol) in dry toluene (2 mL) at room temperature under an argon atmosphere was treated dropwise with trimethylsilyl cyanide (0.32 mL, 2.4 mmol) and the mixture was stirred for 2.5 h. Pyridine (4 mL) and phosphorous oxychloride (0.5 mL, 5.0 mmol) were added, and the solution was refluxed for 40 h. After quenching with ice, the reaction was extracted three times with methylene chloride, the extract was washed with sodium bicarbonate, 10% hydrochloric acid and water, was dried (magnesium sulfate) and the solvent was evaporated. Purification by flash chromatography, eluting with 1:4 ethyl acetate/hexanes,

provided a pale yellow oil (0.3g, 42%). Anal. ($C_{20}H_{22}N_2O_2 \cdot 3/8H_2O$) calcd: C 72.98, H 6.97, N 8.51; found: C 72.85, H 6.80, N 8.56; $R_f = 0.13$ (4:1 hexanes/ethyl acetate).

Example 30

Preparation of 3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1,3-dicarbonitrile

The title compound was isolated as the second product in Example 29 as an off-white solid (0.1g, 19%). mp 77-79°C; Anal. ($C_{20}H_{22}N_2O_2 \cdot 1/4H_2O$) calcd: C 73.48, H 6.94, N 8.57; found: C 73.19, H 6.74, N 8.34; $R_f = 0.32$ (4:1 hexanes/ethyl acetate).

METHODS OF TREATMENT

In order to use a compound of Formula (I) or a pharmaceutically acceptable salt thereof for the treatment of humans and other mammals, it is normally formulated in accordance with standard pharmaceutical practice as a pharmaceutical composition. The compounds of Formula (I) or a pharmaceutically acceptable salt thereof can be used in the manufacture of a medicament for the prophylactic or therapeutic treatment of any disease state in a human or other mammal which is mediated by inhibition of PDE IV, such as but not limited to asthma, allergic, or inflammatory diseases. The compounds of Formula (I) are administered in an amount sufficient to treat such a disease in a human or other mammal.

The method of treatment and monitoring for an HIV-infected human manifesting immune dysfunction or cytokine-mediated disease associated problems is taught in Hanna, WO 90/15534, December 27, 1990. In general, an initial treatment regimen can be copied from that known to be effective in interfering with TNF activity for other TNF mediated disease states by the compounds of Formula (I). Treated individuals will be regularly checked for T cell numbers and T4/T8 ratios and/or measures of viremia such as levels of reverse transcriptase or viral proteins, and/or for progression of monokine-mediated disease associated problems such as cachexia or muscle degeneration. If no effect is seen following the normal treatment regimen, then the amount of the monokine activity interfering agent administered is increased, e.g., by fifty percent per week.

The pharmaceutical composition of the present invention will comprise an effective, non-toxic amount of a compound of Formula (I) and a pharmaceutically acceptable carrier or diluent. The compounds of Formula (I) are administered in conventional dosage forms prepared by combining a compound of Formula (I) in an amount sufficient to produce TNF production inhibiting activity, respectively, with standard pharmaceutical carriers according to conventional procedures. These procedures may involve mixing, granulating, and compressing or dissolving the ingredients as appropriate to the desired preparation.

Thus, if a solid carrier is used, the preparation can be tableted, placed in a hard gelatin capsule in powder or pellet form, or in the form of a troche or lozenge. The amount of solid carrier will vary widely but preferably will be from about 25 mg to about 1 gram.

When a liquid carrier is used, the preparation will be in the form of a syrup, emulsion, soft gelatin capsule, sterile injectable liquid such as an ampule or nonaqueous liquid suspension. Where the composition is in the form of a capsule, any routine encapsulation is suitable, for example using the aforementioned carriers in a hard gelatin capsule shell. Where the composition is in the form of a soft gelatin shell capsule any pharmaceutical carrier routinely used for preparing dispersions or suspensions may be considered, for example aqueous gums, celluloses, silicates, or oils and are incorporated in a soft gelatin capsule shell. A syrup formulation will generally consist of a suspension or solution of the compound or salt in a liquid carrier for example, ethanol, glycerine, or water with a flavoring or coloring agent.

The daily dosage regimen for oral administration is suitably about .001 mg/kg to 100mg/kg, preferably 0.01 mg/Kg to 40 mg/Kg, of a compound of Formula (I) or a pharmaceutically acceptable salt thereof calculated as the free base. The active ingredient may be administered from 1 to 6 times a day, sufficient to exhibit activity.

While it is possible for an active ingredient to be administered neat, it is preferable to present it as a pharmaceutical formulation. The active ingredient may comprise, for topical administration, from 0.001% to 10% w/w, e.g., from 1% to 2% by weight of formulation, although it may comprise as much as 10% w/w but preferably not in excess of 5% w/w and more preferably from 0.1% to 1% w/w of Formulation.

Formulations of the present invention comprise an active ingredient together with one or more acceptable carrier(s) thereof and optionally any other therapeutic ingredient(s). The carrier(s) must be 'acceptable' in the sense of being compatible with the other ingredients of Formulation and not deleterious to the recipient thereof.

It will be recognized by one of skill in the art that the form and character of the pharmaceutically acceptable carrier or diluent is dictated by the amount of active ingredient with which it is to be combined, the route of administration, and other well-known variables.

No toxic effects are expected when these compounds are administered in accordance with the present invention.

Utility Examples

Example A

Inhibitory effect of compounds of Formula (I) on *in vitro* TNF production by human monocytes

The inhibitory effect of compounds of Formula (I) on *in vitro* TNF production by human monocytes may be determined by the protocol as described in Badger *et al.*, EPO published Application 0 411 754 A2, February 6, 1991, and in Hanna, WO 90/15534, December 27, 1990.

Example B

Two models of endotoxic shock have been utilized to determine *in vivo* TNF activity

for the compounds of Formula (I). The protocol used in these models is described in Badger *et al.*, EPO published Application 0 411 754 A2, February 6, 1991, and in Hanna, WO 90/15534, December 27, 1990.

5 The exemplified compounds herein demonstrated a positive *in vivo* response in reducing serum levels of TNF induced by the injection of endotoxin.

Example C

Isolation of PDE Isozymes

10 The phosphodiesterase inhibitory activity and selectivity of the compounds of Formula (I) can be determined using a battery of five distinct PDE isozymes. The tissues used as sources of the different isozymes are as follows: 1) PDE Ib, porcine aorta; 2) PDE Ic, guinea-pig heart; 3) PDE III, guinea-pig heart; 4) PDE IV, human monocyte; and 5) PDE V (also called "Ia"), canine tracheaolis. PDEs Ia, Ib, Ic and III are partially purified using standard chromatographic techniques [Torphy and Cieslinski, Mol. Pharmacol., 37:206-214, 1990]. PDE IV is purified to kinetic homogeneity by the sequential use of
15 anion-exchange followed by heparin-Sepharose chromatography [Torphy *et al.*, J. Biol. Chem., 267:1798-1804, 1992].

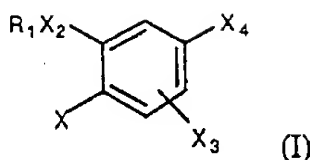
Phosphodiesterase activity is assayed as described in the protocol of Torphy and Cieslinski, Mol. Pharmacol., 37:206-214, 1990. Positive IC₅₀'s in the nanomolar to μ M range for compounds of the workings examples described herein for Formula (I) have been
20 demonstrated.

Example D

The ability of selected PDE IV inhibitors to increase cAMP accumulation in intact tissues is assessed using U-937 cells, a human monocyte cell line that has been shown to contain a large amount of PDE IV. To assess the activity of PDE IV inhibition in intact
25 cells, nondifferentiated U-937 cells (approximately 10^5 cells/reaction tube) were incubated with various concentrations (0.01-1000 μ M) of PDE inhibitors for one minute and 1 μ M prostaglandin E2 for an additional four minutes. Five minutes after initiating the reaction, cells were lysed by the addition of 17.5% perchloric acid, the pH was neutralized by the addition of 1M potassium carbonate and cAMP content was assessed by RIA. A general
30 protocol for this assay is described in Brooker *et al.*, Radioimmunoassay of cyclic AMP and cyclic GMP., Adv. Cyclic Nucleotide Res., 10:1-33, 1979. The compounds of the working examples as described herein for Formula (I) have demonstrated a positive EC₅₀s in the μ M range in the above assay.

What is claimed is:

1. A compound by Formula (I):



wherein:

R_1 is $-(CR_4R_5)_nC(O)O(CR_4R_5)_mR_6$, $-(CR_4R_5)_nC(O)NR_4(CR_4R_5)_mR_6$, $-(CR_4R_5)_nO(CR_4R_5)_mR_6$, or $-(CR_4R_5)_rR_6$ wherein the alkyl moieties may be optionally substituted with one or more halogens;

m is 0 to 2;

n is 1 to 4;

r is 0 to 6;

R_4 and R_5 are independently selected from hydrogen or a C₁₋₂ alkyl;

R_6 is hydrogen, methyl, hydroxyl, aryl, halo substituted aryl, aryloxyC₁₋₃ alkyl, halo substituted aryloxyC₁₋₃ alkyl, indanyl, indenyl, C₇₋₁₁ polycycloalkyl, tetrahydrofuranyl, furanyl, tetrahydropyranyl, pyranal, tetrahydrothienyl, thienyl, tetrahydrothiopyranal, thiopyranal, C₃₋₆ cycloalkyl, or a C₄₋₆ cycloalkyl containing one or two unsaturated bonds, wherein the cycloalkyl and heterocyclic moieties may be optionally substituted by 1 to 3 methyl groups or one ethyl group;

provided that:

a) when R_6 is hydroxyl, then m is 2; or

b) when R_6 is hydroxyl, then r is 2 to 6; or

c) when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranal, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then m is 1 or 2; or

d) when R_6 is 2-tetrahydropyranyl, 2-tetrahydrothiopyranal, 2-tetrahydrofuranyl, or 2-tetrahydrothienyl, then r is 1 to 6;

e) when n is 1 and m is 0, then R_6 is other than H in $-(CR_4R_5)_nO(CR_4R_5)_mR_6$;

X is YR_2 , halogen, nitro, NR_4R_5 , or formyl amine;

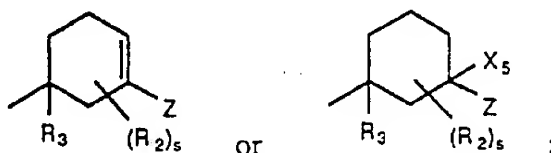
Y is O or $S(O)m'$;

m' is 0, 1, or 2;

X_2 is O or NR_8 ;

X_3 is hydrogen or X;

X_4 is



(a)

(b)

X₅ is H, R₉, OR₈, CN, C(O)R₈, C(O)OR₈, C(O)NR₈R₈, or NR₈R₈;

R₂ is independently -CH₃ or -CH₂CH₃ optionally substituted by 1 or more halogens;

5 s is 0 to 4;

R₃ is hydrogen, halogen, C₁₋₄ alkyl, CH₂NHC(O)C(O)NH₂, halo-substituted C₁₋₄ alkyl, -CH=CR₈R₈', cyclopropyl optionally substituted by R₈', CN, OR₈, CH₂OR₈, NR₈R₁₀, CH₂NR₈R₁₀, C(Z')H, C(O)OR₈, C(O)NR₈R₁₀, or C≡CR₈;

10 Z' is O, NR₉, NOR₈, NCN, C(-CN)₂, CR₈CN, CR₈NO₂, CR₈C(O)OR₈, CR₈C(O)NR₈R₈, C(-CN)NO₂, C(-CN)C(O)OR₉, or C(-CN)C(O)NR₈R₈;

Z is C(Y')R₁₄, C(O)OR₁₄, C(Y')NR₁₀R₁₄, C(NR₁₀)NR₁₀R₁₄, CN, C(NOR₈)R₁₄, C(O)NR₈NR₈C(O)R₈, C(O)NR₈NR₁₀R₁₄, C(NOR₁₄)R₈, C(NR₈)NR₁₀R₁₄, C(NR₁₄)NR₈R₈, C(NCN)NR₁₀R₁₄, C(NCN)SR₉, (2-, 4- or 5-imidazolyl), (3-, 4- or 5-pyrazolyl), (4- or 5-triazolyl[1,2,3]), (3- or 5-triazolyl[1,2,4]), (5-tetrazolyl), (2-, 4- or 5-oxazolyl), (3-, 4- or 5-isoxazolyl), (3- or 5-oxadiazolyl[1,2,4]), (2-oxadiazolyl[1,3,4]), (2-thiadiazolyl[1,3,4]), (2-, 4-, or 5-thiazolyl), (2-, 4-, or 5-oxazolidinyl), (2-, 4-, or 5-thiazolidinyl), or (2-, 4-, or 5-imidazolidinyl); wherein all of the heterocyclic ring systems may be optionally substituted one or more times by R₁₄;

Y' is O or S;

20 R₇ is -(CR₄R₅)_qR₁₂ or C₁₋₆ alkyl wherein the R₁₂ or C₁₋₆ alkyl group is optionally substituted one or more times by C₁₋₂ alkyl optionally substituted by one to three fluorines, -F, -Br, -Cl, -NO₂, -NR₁₀R₁₁, -C(O)R₈, -C(O)OR₈, -OR₈, -CN, -C(O)NR₁₀R₁₁, -OC(O)NR₁₀R₁₁, -OC(O)R₈, -NR₁₀C(O)NR₁₀R₁₁, -NR₁₀C(O)R₁₁, -NR₁₀C(O)OR₉, -NR₁₀C(O)R₁₃, -C(NR₁₀)NR₁₀R₁₁, -C(NCN)NR₁₀R₁₁, -C(NCN)SR₉, -NR₁₀C(NCN)SR₉, -NR₁₀C(NCN)NR₁₀R₁₁, -NR₁₀S(O)₂R₉, -S(O)_mR₉, -NR₁₀C(O)C(O)NR₁₀R₁₁, -NR₁₀C(O)C(O)R₁₀, thiazolyl, imidazolyl, oxazolyl, pyrazolyl, triazolyl, or tetrazolyl;

q is 0, 1, or 2;

30 R₁₂ is C₃₋₇ cycloalkyl, (2-, 3- or 4-pyridyl), pyrimidyl, pyrazolyl, (1- or 2-imidazolyl), thiazolyl, triazolyl, pyrrolyl, piperazinyl, piperidinyl, morpholinyl, furanyl, (2- or 3-thienyl), (4- or 5-thiazolyl), quinolinyl, naphthyl, or phenyl;

R₈ is hydrogen or R₉;

R₈' is R₈ or fluorine;

R₉ is C₁₋₄ alkyl optionally substituted by one to three fluorines;

35 R₁₀ is OR₈ or R₁₁;

R₁₁ is hydrogen, or C₁₋₄ alkyl optionally substituted by one to three fluorines; or when R₁₀ and R₁₁ are as NR₁₀R₁₁ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom which is O, N, or S;

R₁₃ is oxazolidinyl, oxazolyl, thiazolyl, pyrazolyl, triazolyl, tetrazolyl, imidazolyl, imidazolidinyl, thiazolidinyl, isoxazolyl, oxadiazolyl, or thiadiazolyl, and each of these heterocyclic rings is connected through a carbon atom and each may be unsubstituted or substituted by one or two C₁₋₂ alkyl groups;

5 R₁₄ is hydrogen or R₇; or when R₁₀ and R₁₄ are as NR₁₀R₁₄ they may together with the nitrogen form a 5 to 7 membered ring optionally containing at least one additional heteroatom which is O, N, or S;

provided that:

10 f) when R₁₂ is N-pyrazolyl, N-imidazolyl, N-triazolyl, N-pyrrolyl, N-piperazinyl, N-piperidinyl, or N-morpholinyl, then q is not 1; or
or the pharmaceutically acceptable salts thereof.

2. A compound of claim 1 which is :

methyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate; SB 212179

15 ethyl 5-cyano-5-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate; SB

methyl 5-cyano-5-(3-cyclopentylloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate; SB 211572

20 methyl 5-cyano-5-(3,4-bisdifluoromethoxyphenyl)cyclohex-1-ene-1-carboxylate; SB

methyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate]; SB 210667

ethyl *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate]; SB 211600

25 methyl *cis*-[3-cyano-3-(3-cyclopentylloxy-4-methoxyphenyl)cyclohexane-1-carboxylate]; SB 212066

methyl *cis*-[3-(3,4-bisdifluoromethoxyphenyl)-3-cyanocyclohexane-1-carboxylate]; SB

30 *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid]; SB 210984

cis-[3-cyano-3-(3-cyclopentylloxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid]; SB 212510

cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide]; SB 211529

35 *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-N-methylcarboxamide]; SB 213021

cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-N,N-dimethylcarboxamide]; SB 212697

cis-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-(N-(4-bromobenzyl)carboxamide)]; SB 212698

cis-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxamide]; SB 214236

5 *cis*-[3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1,3-dicyano-cyclohexane]; SB 212188

cis-[3-(3-cyclopentyloxy-4-methoxyphenyl)-1,3-dicyanocyclohexane]; SB 213832

10 *cis*-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1-(3-methyl[1,2,4]oxadiazol-5-yl)cyclohexane]; SB 213826

cis-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)-1-(3-methyl[1,2,4]-oxadiazol-5-yl)cyclohexane]; SB 214243

trans-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)-cyclohexane-1-carboxylate]; SB 213677

15 *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)-cyclohexane-1-carboxylate];, SB 213951

trans-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid];, SB 213731

20 *trans*-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxylic acid];, SB 213921

trans-[3-cyano-3-(3-cyclopropylmethoxy-4-methoxyphenyl)cyclohexane-1-carboxamide]; SB 213835

trans-[3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohexane-1-carboxamide]; SB 213921

25 *trans*-[3-(3-cyclopropylmethoxy-4-methoxyphenyl)-1,3-dicyano-cyclohexane]; SB 213920

trans-[3-(3-cyclopentyloxy-4-methoxyphenyl)-1,3-dicyanocyclohexane]; and SB 214241

30 *trans*-[3-cyano-3-(3-cyclopentyloxyphenyl)-1-(3-methyl[1,2,4]oxadiazol-5-yl)cyclo-hexane]; SB 214242

methyl 3-cyano-3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate; methyl 5-cyano-5-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1-carboxylate;

35 3-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1,3-dicarbonitrile; or 5-(3-cyclopentyloxy-4-methoxyphenyl)cyclohex-1-ene-1,5-dicarbonitrile.

3. A pharmaceutical composition comprising a compound of Formula (I) according to claim 1 and a pharmaceutically acceptable excipient.

4. A method for treating an allergic or inflammatory state which method

comprises administering to a subject in need thereof an effective amount of a compound of Formula (I) according to claim 1 alone or in combination with a pharmaceutically acceptable excipient.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US94/08581

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61K 31/275, 31/42; C07C 255/45, 271/06

US CL : 514/364, 521, 523, 525; 548/131, 426

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/364, 521, 523, 525; 548/131, 426

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS ONLINE SEARCH OF STRUCTURE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US, A, 2,573,015 (HAFLIGER ET AL) 30 OCTOBER 1951, SEE COLUMN 3.	1-4
A	US, A, 2,650,243 (NOVELLO) 25 AUGUST 1953, SEE EXAMPLES.	1-4
A	US, A, 2,822,383 (SMITH) 04 FEBRUARY 1958, SEE EXAMPLE 1.	1-4
A	US, A, 2,886,589 (NOVELLO) 12 MAY 1959, SEE EXAMPLES.	1-4
A	US, A, 2,911,432 (HOEHM) 03 NOVEMBER 1959, SEE FORMULA	1-4

☐ Further documents are listed in the continuation of Box C.
 ☐ See patent family annex.

* Special categories of cited documents:	* T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
* A* document defining the general state of the art which is not considered to be of particular relevance	* X	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
* E* earlier document published on or after the international filing date	* Y	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
* L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	* Z	document member of the same patent family
* O* document referring to an oral disclosure, use, exhibition or other means		
* P* document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

28 OCTOBER 1994

Date of mailing of the international search report

NOV 07 1994

 Name and mailing address of the ISA/US
 Commissioner of Patents and Trademarks
 Box PCT
 Washington, D.C. 20231

 Authorized officer
 JOSEPH BRUST

Facsimile No. (703) 305-3230

Telephone No. (703) 308-1235

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/08581**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☒ Claims Nos.: 1,3,4 (in part)
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

Please See Extra Sheet.

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

BOX 1. OBSERVATIONS WHERE CLAIMS WERE FOUND UNSEARCHABLE

2. Where no meaningful search could be carried out, specifically:

Generic claim 1, is such an "alphabet soup" of variables with such a multitude and plethora of variables that no meaningful specific compounds can be deduced therefrom save those specifically defined on page 12, line 5 to page 13, line 2, and in the Examples 1-30, and in claim 2. Hence, the above claims will only be searched to the extent that they read on those above noted searchable areas.